In the Claims:

Please cancel claims 2-18 without prejudice.

Please amend Claim 1 and add new claims 19-31 as follows:

1. (once amended) A process for synthesizing a dihydroindole C-ring of a CC-1065/duocarmycin analog, the dihydroindole C-ring of a CC-1065/duocarmycin analog being represented by the following structure:

the process comprising the following steps:

Step A: allylating an ortho-haloaniline with 1,3-dichloropropene for forming a vinyl chloride, the *ortho*-haloaniline being represented by the following structure:

$$R^2$$
 $R^3$ 
 $R^4$ 
 $NH$ 
 $BOC$ 

wherein:

R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, and R<sup>4</sup> are radicals independently selected from the group consisting of hydrogen, alkyl(C1-C6), alkoxy, and arylalkoxy, with a proviso that R<sup>1</sup> and R<sup>2</sup>, or R<sup>2</sup> and R<sup>3</sup> may form a fused 5- or 6-membered ring with or without a heteroatom; and

X is a halide selected from the group consisting of bromine and iodine; and the vinyl chloride is represented by the following structure:

$$R^2$$
 $R^3$ 
 $R^4$ 
 $R^4$ 

Page -2-



Step B: cyclizing the vinyl chloride of said step A for forming the dihydroindole C-ring of the CC-1065 / duocarmycin analog.

A3

- 19. (new claim) A process according to claim 1 wherein, in said Step A, the *ortho*-haloaniline is an *ortho*-bromoaniline.
- 20. (new claim) A process according to claim 1 wherein, in said Step A, the *ortho*-haloaniline is an *ortho*-iodoaniline.
- 21. (new claim) A process according to claim 1 wherein, in said Step A, the *ortho*-haloaniline is protected with a BOC group.
- 22. (new claim) A process according to claim 1 wherein, in said Step A, said allylation is catalyzed by the addition of a catalytic amount of tetra-*n*-butyl ammonium iodide.
- 23. (new claim) A process according to claim 1 wherein, in said Step B, said cyclization is performed with an addition of tri-*n*-butyltin hydride.
- 24. (new claim) A process according to claim 23 wherein, in said Step B, said cyclization is catalyzed by the addition of a catalytic amount of AIBN.
- 25. (new claim) A process according to claim 24 wherein, in said Step B, said cyclization is performed using toluene as the solvent.
- 26. (new claim) A process according to claim 1 wherein: in said Step A, the vinyl chloride is represented by the following structure:

Page -3-

in said Step B, the dihydroindole C-ring of the CC-1065 / duocarmycin analog is represented by the following structure:

27. (new claim) A process according to claim 1 wherein:

in said Step A, the vinyl chloride is represented by the following structure:

in said Step B, the dihydroindole C-ring of the CC-1065 / duocarmycin analog is represented by the following structure:

28. (new claim) A process according to claim 1 wherein:

in said Step A, the vinyl chloride is represented by the following structure:

in said Step B, the dihydroindole C-ring of the CC-1065 / duocarmycin analog is represented by the following structure:

29. (new claim) A process according to claim 1 wherein:

in said Step A, the vinyl chloride is represented by the following structure:

in said Step B, the dihydroindole C-ring of the CC-1065 / duocarmycin analog is represented by the following structure:

30. (new claim) A process according to claim 1 wherein:

in said Step A, the vinyl chloride is represented by the following structure:

in said Step B, the dihydroindole C-ring of the CC-1065 / duocarmycin analog is represented by the following structure:

Page -5-

31. (new claim) A process according to claim 1 wherein:

in said Step A, the vinyl chloride is represented by the following structure:

in said Step B, the dihydroindole C-ring of the CC-1065 / duocarmycin analog is represented by the following structure:

Respectfully submitted,

O. f 30, 2001

Date

Donald G. Lewis, Reg. No. 28,636

THE SCRIPPS RESEARCH INSTITUTE

Office of Patent Counsel

10550 North Torrey Pines Road

Mail Drop: TPC-8

La Jolla, California 92037

October 30, 2001

(858) 784-2937